## **AMENDMENTS TO THE CLAIMS:**

Please amend the claims as follows:

- 1. (Currently amended) A promoter comprising an [[An]] isolated DNA molecule comprising a nucleotide sequence having a nucleotide sequence of at least 300 nucleotides, wherein the selected from the group consisting of:
- [[a) a]] nucleotide sequence [[that]] has at least [[90%]] <u>95%</u> sequence identity to the sequence set forth in SEQ ID NO:8[[; and]] or the
- [[b) a]] nucleotide sequence [[that]] hybridizes to the sequence set forth in SEQ ID NO:8 under at least high stringency conditions.

Claims 2-69. (Cancelled).

- 70. (Currently amended) A promoter comprising an [[An]] isolated DNA molecule comprising a nucleotide sequence having a nucleotide sequence of at least 300 nucleotides, wherein the nucleotide sequence [[that]] has at least [[90%]] 95% sequence identity to the sequence set forth in SEQ ID NO:8 or the nucleotide sequence hybridizes to the sequence set forth in SEQ ID NO:8 under at least high stringency conditions, wherein the nucleotide sequence is further selected from the group consisting of:
  - a) a nucleotide sequence set forth in SEQ ID NO:6;
- b) a nucleotide sequence that has at least **[[90%]]** <u>95%</u> sequence identity to the sequence set forth in SEQ ID NO:6;
- c) a nucleotide sequence that hybridizes to the sequence set forth in SEQ ID NO:6 under at least high stringency conditions;
  - d) the nucleotide sequence set forth in SEQ ID NO:7;
- e) a nucleotide sequence that has at least **[[90%]]** <u>95%</u> sequence identity to the sequence set forth in SEQ ID NO:7; and
- f) a nucleotide sequence that hybridizes to the sequence set forth in SEQ ID NO:7 under **at least** high stringency conditions.
- 71. (Currently amended) The DNA-molecule promoter of claim 1, wherein the

nucleotide sequence <u>as set forth in SEQ ID NO:6 or 7</u> is <del>obtained</del> <u>derived</u> from a virus.

- 72. **(Currently amended)** The **DNA molecule promoter** of claim 1, wherein the nucleotide sequence **as set forth in SEQ ID NO:6 or 7** is **obtained derived** from a badnavirus.
- 73. (Currently amended) A monocotyledonous plant comprising a nucleotide sequence according to The DNA molecule of claim 71, wherein a nucleic acid operably linked to the nucleotide sequence as set forth in SEQ ID NO:6 or 7 is expressed constitutively in [[a]] the monocotyledonous plant.
- 74. (Currently amended) A non-graminaceous monocotyledonous plant comprising a nucleotide sequence according to The DNA molecule of claim 71, wherein a nucleic acid operably linked to the nucleotide sequence as set forth in SEQ ID NO:6 or 7 is expressed constitutively in [[a]] the non-graminaceous monocotyledonous plant.
- 75. (Currently amended) The <u>non-graminaceous monocotyledonous plant</u> DNA molecule of claim 74, wherein the non-graminaceous monocotyledonous plant is selected from the group consisting of *Musaceae*, taro, ginger, onions, garlic, pineapple, bromeliaeds, palms, orchids, lilies and irises.
- 76. (Currently amended) The <u>non-graminaceous monocotyledonous plant</u> <del>DNA</del> molecule of claim 74, wherein the non-graminaceous monocotyledonous plant is taro.
- 77. **(Currently amended)** The **DNA molecule promoter** of claim 1, wherein **[[the]]** the nucleotide sequence is as set forth in SEQ ID NO:6 or 7.
- 78-80. (Cancelled).
- 81. (Withdrawn) An isolated polynucleotide comprising a nucleotide sequence that corresponds or is complementary to at least a portion of the sequence set forth in SEQ ID NO:1 or to a variant thereof wherein the portion is at least 90 nucleotides in length and wherein the variant displays at least 80% sequence identity to the at least a portion.

- 82. (Withdrawn) polynucleotide of claim 81, wherein the variant displays at least 85% sequence identity to at the least a portion.
- 83. (Withdrawn) The polynucleotide of claim 82, wherein the variant displays at least 80% sequence identity to a nucleotide sequence that encodes an amino acid sequence selected from the group consisting of SEQ ID NO:3, 4 and 5.
- 84. (Withdrawn) The polynucleotide of claim 81, wherein the variant hybridises to at least a portion of the sequence set forth in SEQ ID NO:1, which is at least 18 nucleotides in length, under at least high stringency conditions.
- 85. (Withdrawn) The polynucleotide of claim 84, wherein the variant hybridises to a nucleotide sequence that encodes an amino acid sequence selected from the group consisting of SEQ ID NO:3, 4 and 5 under at least high stringency conditions.
- 86. (Withdrawn) An isolated polypeptide comprising an amino acid sequence selected from the group consisting of:
- (i) at least a portion of the sequence set forth in SEQ ID NO:4, wherein the portion is at least six amino acids in length;
- (ii) at least a portion of a variant that displays at least 55% sequence identity to the sequence set forth in SEQ ID NO:4, wherein the portion is at least 15 amino acid residues in length;
- (iii) at least a portion of the sequence set forth in SEQ ID NO:5, wherein the portion is at least seven amino acids in length;
- (iv) at least a portion of a variant that displays at least 65% sequence identity to the sequence set forth in SEQ ID NO:5, wherein the portion is at least 30 amino acid residues in length:
- (v) least a portion of the sequence set forth in SEQ ID NO:6, wherein the portion is at least 16 amino acid residues in length;
- (vi) at least a portion of a variant that displays at least 70% sequence identity to the sequence set forth in SEQ ID NO:6, wherein the portion is at least 30 amino acid residues in length.

- 87. (Currently amended) A chimeric DNA construct comprising the nucleotide sequence of claim 1, as set forth in SEQ ID NO:6 or 7, operably linked to a foreign or endogenous DNA sequence to be transcribed.
- 88. (Currently amended) The construct of claim 87, further comprising a 3' non-translated sequence that is operably linked to the **foreign or endogenous** DNA sequence to be transcribed and that functions in plant cells to terminate transcription and/or to cause addition of a polyadenylated nucleotide sequence to the 3' end of a transcribed RNA sequence.
- 89. (Currently amended) A chimeric DNA construct comprising the nucleotide sequence of claim 70, wherein the nucleotide sequence as set forth in SEQ ID NO:6 or 7 is operably linked to a foreign or endogenous DNA sequence to be transcribed.
- 90. (Cancelled).
- 91. (Currently amended) The construct of claim 87, wherein the nucleotide sequence <a href="https://example.com/hybridizes to the sequence">hybridizes to the sequence</a> [[is]] as set forth in SEQ ID NO:6 or 7 <a href="https://example.com/under-thigh-stringency-conditions">under thigh stringency conditions</a>.
- 92. (Cancelled).
- 93. **(Currently amended)** The construct of claim 87, wherein the **foreign or endogenous** DNA sequence **to be transcribed** encodes a structural or regulatory protein.
- 94. **(Currently amended)** The construct of claim 87, wherein the **foreign or endogenous** DNA sequence **to be transcribed** encodes a transcript capable of modulating expression of a corresponding target gene.
- 95. **(Currently amended)** The construct of claim 94, wherein the transcript comprises a transcribed region **aimed at <u>for</u>** downregulating the expression of the corresponding target gene.
- 96. (Currently amended) The construct of claim 94, wherein the transcript comprises a transcribed region comprising that represents a molecules selected from the group

- consisting of a sense suppression molecule, an antisense RNA, a ribozyme and an RNAi molecule.
- 97. (Previously presented) The construct of claim 87, further comprising an enhancer element.
- 98. (Previously presented) The construct of claim 87, further comprising a leader sequence which modulates mRNA stability.
- 99. (Currently amended) The construct of claim 87, further comprising <u>a nucleic acid</u> <u>sequence encoding</u> a targeting sequence for targeting a protein product of the <u>foreign</u> <u>or endogenous</u> DNA sequence <u>to be transcribed</u> to an intracellular compartment within plant cells or to an extracellular environment.
- 100. (Previously presented) The construct of claim 87, further comprising a selectable marker gene.
- 101. (Previously presented) The construct of claim 87, further comprising a screenable marker gene.
- 102. (Currently amended) A host cell transformed with a [[The]] construct of claim 87, wherein a nucleic acid operably linked to the promoter or biologically active fragment or variant is constitutively expressed in a host cell.
- 103. (Currently amended) The <u>host cell</u> construct of claim 102, wherein the host cell is a plant cell.
- 104. (Currently amended) The <u>host cell</u> construct of claim 102, wherein the host cell is a monocotyledonous plant cell.
- 105. (Currently amended) The <u>host cell</u> construct of claim 102, wherein the host cell is a non-graminaceous monocotyledonous plant cell.
- 106. (Currently amended) The <u>host cell</u> construct of claim 102, wherein the host cell is a non-graminaceous monocotyledonous plant cell selected from the group consisting of *Musaceae*, taro, ginger, onions, garlic, pineapple, bromeliaeds, palms, orchids, lilies and irises.

- 107. **(Currently amended)** The <u>host cell</u> <del>construct</del> of claim 102<del>, wherein the host cell</del> is a graminaceous monocotyledonous plant cell.
- 108. (Currently amended) The <u>host cell</u> construct of claim 102, wherein the host cell is a dicotyledonous plant cell.
- 109. (Withdrawn) A method for gene expression in a plant, comprising introducing into a plant cell a chimeric DNA construct comprising an isolated promoter or biologically active fragment thereof or variant of these, wherein the promoter is naturally located upstream of a transcribable DNA sequence which hybridises to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein the promoter or biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed.
- 110. (Withdrawn) A method for producing transformed plant cells, comprising:
- (a) introducing into regenerable plant cells a chimeric DNA construct comprising an isolated promoter or biologically active fragment thereof or variant of these, wherein the promoter is naturally located upstream of a transcribable DNA sequence which hybridises to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein the promoter or biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed, so as to yield transformed plant cells; and
  - (b) identifying or selecting transformed plant cells.
- 111. (Withdrawn) A method for selecting stable genetic transformants from transformed plant cells comprising:
- (a) introducing into regenerable plant cells a chimeric DNA construct comprising an isolated promoter or biologically active fragment thereof or variant of these, wherein the promoter is naturally located upstream of a transcribable DNA sequence which hybridises to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein the promoter or

biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed, so as to yield transformed plant cells; and

- (b) identifying or selecting a transformed plant cell line from said transformed plant cells.
- 112. (Withdrawn) A method for producing a differentiated transgenic plant, comprising:
- (a) introducing into regenerable plant cells a chimeric DNA construct comprising an isolated promoter or biologically active fragment thereof or variant of these, wherein the promoter is naturally located upstream of a transcribable DNA sequence which hybridises to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein the promoter or biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed, so as to yield regenerable transformed plant cells;
  - (b) identifying or selecting a population of transformed plant cells; and
  - (c) regenerating a differentiated transgenic plant from the population.
- 113. (Withdrawn) The method of claims 109, wherein the cells are dicotyledonous plant cells.
- 114. (Withdrawn) The method claim 109, wherein the cells are monocotyledonous plant cells.
- 115. (Withdrawn) The method of claim 109, wherein the cells are graminaceous monocotyledonous plant cells.
- 116. (Withdrawn) The method of claim 109, wherein the cells are non-graminaceous monocotyledonous plant cells.
- 117. (Withdrawn) The method of claim 109, wherein expression of the chimeric DNA construct in the transformed cells imparts a phenotypic characteristic to the transformed cells.
- 118. (Withdrawn) The method of claim 109, wherein the construct comprises a selectable marker gene.

- 119. (Withdrawn) The method of claim 109, wherein the construct comprises a screenable marker gene.
- 120. (Withdrawn) The method of claim 112, wherein expression of the chimeric DNA construct renders the differentiated transgenic plant identifiable over the corresponding non-transgenic plant.
- 121. (Withdrawn) The method of claim 112, further comprising obtaining progeny from the differentiated transgenic plant.
- 122. (Withdrawn) Progeny obtained by the method of claim 121.
- 123. (Withdrawn) A plant part of the differentiated transgenic plant obtained from the method of claim 112, wherein the plant part contains the chimeric construct.
- 124. (Withdrawn) A differentiated transgenic plant regenerated from transformed plant cells obtained by the method of claim 110.
- 125. (Withdrawn) A transformed plant cell containing a chimeric DNA construct comprising an isolated plant promoter or biologically active fragment thereof or variant of these, wherein said promoter is naturally located upstream of a transcribable DNA sequence which hybridises to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein said promoter or biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed.
- 126. (Withdrawn) A differentiated transgenic plant comprising plant cells containing a chimeric DNA construct comprising an isolated plant promoter or biologically active fragment thereof or variant of these, wherein said promoter is naturally located upstream of a transcribable DNA sequence which hybridises to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein said promoter or biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed.
- 127. (Withdrawn) The transgenic plant of claim 126, wherein the plant is a dicotyledonous plant.

- 128. (Withdrawn) The transgenic plant of claim 126, wherein the plant is a monocotyledonous plant.
- 129. (Withdrawn) The transgenic plant of claim 126, wherein the plant is a graminaceous monocotyledonous plant.
- 130. (Withdrawn) The transgenic plant of claim 126, wherein the plant is a non-graminaceous monocotyledonous plant.
- 131. (Withdrawn) The transgenic plant of claim 126, wherein the construct comprises a selectable marker gene.
- 132. (Withdrawn) The transgenic plant of claim 126, wherein the construct comprises a screenable marker gene.
- 133. (Withdrawn) The transgenic plant of claim 126, wherein the expression of the chimeric DNA construct renders the differentiated transgenic plant identifiable over the corresponding non-transgenic plant.
- 134. (Withdrawn) A method of using of a chimeric DNA construct comprising an isolated plant promoter or biologically active fragment thereof or variant of these, wherein said promoter is naturally located upstream of a transcribable DNA sequence which hybridizes to a nucleic acid probe derived from the polynucleotide sequence set forth in SEQ ID NO:1 under at least high stringency conditions, wherein said promoter or biologically active fragment or variant is operably linked to a foreign or endogenous DNA sequence to be transcribed, in the production of a transformed plant cell, plant or plant part.
- 135. (Withdrawn) A method for diagnosing a badnaviral infection of a plant, comprising detecting the presence in a cell or tissue of the plant of (a) a nucleotide sequence that corresponds or is complementary to at least a portion of the nucleotide sequence set forth in SEQ ID NO:1 or 2, or of a variant of the nucleotide sequence, or (b) an amino acid sequence that corresponds to at least a portion of the sequence set forth in SEQ ID NO:3, 4 or 5, or of a variant of the amino acid sequence.
- 136. (Withdrawn) A method of screening for an agent that modulates badnaviral infection, the method comprising:

- contacting a preparation comprising:
- (i) a polypeptide comprising an amino acid sequence that corresponds to at least a portion of the sequence set forth in SEQ ID NO: 3, 4 or 5, or of a variant of the sequence; or
- (ii) a polynucleotide comprising a nucleotide sequence that corresponds or is complementary to at least a portion of the sequence set forth in SEQ ID NO:1 or 2, which polynucleotide is operably linked to a promoter; or
- (iii) a polynucleotide comprising a reporter gene that is operably connected to a promoter comprising the sequence set forth in SEQ ID NO:6, 7, 8 or 9, with a test agent; and
- detecting a change in the level and/or functional activity of the polypeptide, or an expression product of the nucleotide sequence or of the reporter gene, relative to a normal or reference level and/or functional activity in the absence of the test agent.
- 137. (Withdrawn) The method of claim 136, wherein the agent inhibits or reduces badnavirus infection and the method comprises detecting a reduction in the level and/or functional activity of the polypeptide, or an expression product of the nucleotide sequence or of the reporter gene, relative to the normal or reference level and/or functional activity.
- 138. (Withdrawn) A method for treating and/or preventing badnaviral infection of a plant, comprising administering to the plant an agent that:
  - reduces the level and/or functional activity of:

a polypeptide that comprises an amino acid sequence corresponding to at least a portion of the sequence set forth in SEQ ID NO: 3, 4 or 5, or of a variant of the sequence; or

an expression product of a nucleotide sequence that corresponds or is complementary to at least a portion of the sequence set forth in SEQ ID NO:1 or 2; or

 reduces the functional activity of a promoter that comprises the sequence set forth in any one of SEQ ID NO:6, 7, 8 or 9.